L	Hits	Search Text	DB	Time stamp
Number				_
1 ,	67	cox\$.in. and mismatch	USPAT;	2003/08/26
			US-PGPUB	17:27
2	3	cox\$.in. and mismatch.ti.	USPAT;	2003/08/26
			US-PGPUB	17:28
3	1	mismatch near3 (corepair or co adj1	USPAT;	2003/08/26
		repair)	US-PGPUB	17:29

## (FILE 'HOME' ENTERED AT 17:32:59 ON 26 AUG 2003)

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FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 17:33:06 ON 26 AUG 2003
              9 S MISMATCH? (5A) (COREPAIR OR CO(W)REPAIR)
L1
              3 DUP REM L1 (6 DUPLICATES REMOVED)
L2
     FILE 'SCISEARCH' ENTERED AT 17:34:00 ON 26 AUG 2003
           5411 S CARRAWAY?/RAU
L3
T<sub>1</sub>4
           2307 S L3 AND 1993/RPY
            180 S L4 AND 175/RVL
L5
              0 S L5 AND MULTIPLEX?
L6
             16 S L5 AND MULTIPL?
L7
=> d 15 126 bib ab
     ANSWER 126 OF 180 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN
T.5
     96:94649 SCISEARCH
AN
     The Genuine Article (R) Number: TQ968
GA
     A NOVEL IN-VIVO METHOD TO DETECT DNA-SEQUENCE VARIATION
TI
ΑU
     FAHAM M; COX D R (Reprint)
     STANFORD UNIV, SCH MED, DEPT GENET & PEDIAT, STANFORD, CA, 94305
CS
     (Reprint); STANFORD UNIV, SCH MED, DEPT GENET & PEDIAT, STANFORD, CA,
     94305; UNIV CALIF SAN FRANCISCO, DEPT BIOCHEM & BIOPHYS, SAN FRANCISCO,
     CA, 94143
CYA
     GENOME RESEARCH, (DEC 1995) Vol. 5, No. 5, pp. 474-482.
     ISSN: 1054-9803.
DT
     Article; Journal
     LIFE
FS
     ENGLISH
LА
REC
    Reference Count: 27
     *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*
        Mismatch repair detection (MRD) is an in vivo method that uses a change
AB
     in bacterial colony color to detect DNA sequence variation. DNA fragments
     to be screened for variation are cloned into two MRD plasmids, and
     bacteria are transformed with heteroduplexes of these constructs. The
     resulting colonies are blue in the absence of a mismatch and white in the
     presence of a mismatch. MRD is capable of detecting a single mismatch in a
     DNA fragment as large as 10kb in size. In addition, MRD has the potential
     for analyzing many fragments simultaneously, offering a powerful method
     for high-throughput genotyping and mutation detection in a large genomic
     region.
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